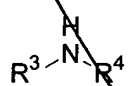
in which

R^1, R^2, R^5 and R^6 have the meanings indicated above,

and then with thionyl chloride and the product thus obtained is reacted in situ in an inert solvent with an amine of the formula (IV)



(IV),

in which

R^3 and R^4 have the meaning indicated above,

and, if appropriate, reacted to give the corresponding salts, hydrates or N-oxides.

Remarks / Explanations

As a result of this preliminary amendment, claims 1-5 remain pending in the application. No new matter has been added.

Claim 1 has been amended in structural formula (I) to show the substitutents on the left-most ring more correctly, and in structural formula (II) to move the group R^5 away from the ring for clarity.

Attached hereto is a marked-up version of the changes made to the specification and claims by the current amendment. The attached page is captioned **"Version with markings to show changes made."**

In view of the above amendments and explanations, this application is deemed to be in condition for allowance, and allowance is accordingly requested.

Respectfully submitted,

William F. Gray

Reg. No. 31018
Phone: (203) 812-2712
Date: 17 Dec '01

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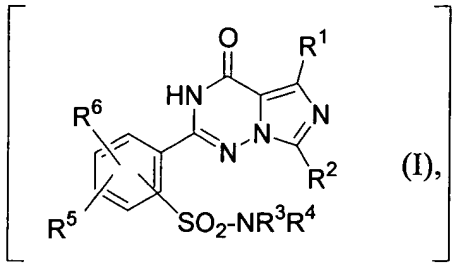
Bayer Corporation
400 Morgan Lane
West Haven, CT 06516-4175

Version with markings to show changes made:

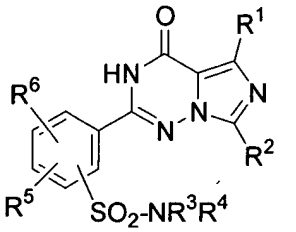
In the claims:

Claim 1 has been amended as shown below:

1. (Amended) Process for the preparation of compounds of the formula I



(I),



(I),

in which

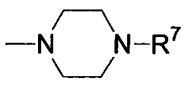
R^1 represents hydrogen or straight-chain or branched alkyl having up to 4 carbon atoms,

R^2 represents straight-chain or branched alkyl having up to 4 carbon atoms,

R^3 and R^4 are identical or different and represent a straight chain or branched alkyl chain having up to 5 carbon atoms, which is optionally substituted up to two times in an identical or different manner by hydroxyl or methoxy,

or

R^3 and R^4, together with the nitrogen atom, form a piperidinyl, morpholinyl or thiomorpholinyl ring or a radical of the formula



in which

R^7 denotes hydrogen, formyl, straight-chain or branched acyl or alkoxycarbonyl each having up to 6 carbon atoms, or straight-chain or branched alkyl having up to 6 carbon atoms, which is optionally mono- to disubstituted, in an identical or different manner, by hydroxyl, carboxyl, straight-chain or branched alkoxy or alkoxycarbonyl each having up to 6 carbon atoms, or denotes C_{3-8} -cycloalkyl,

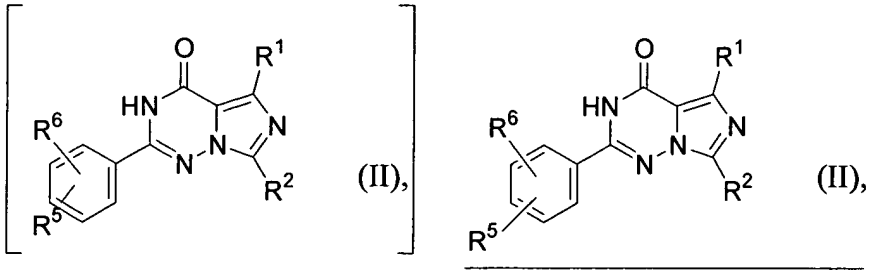
and the heterocycles mentioned under R^3 and R^4, formed together with the nitrogen atom, are optionally mono-to disubstituted, in an identical or different manner, if appropriate also geminally, by hydroxyl, formyl, carboxyl, straight-chain or branched acyl or alkoxycarbonyl each having up to 6 carbon atoms,

and/or the heterocycycles mentioned under R^3 and R^4, formed together with the nitrogen atom, are optionally substituted by straight-chain or branched alkyl having up to 6 carbon atoms, which is optionally mono- to disubstituted, in an identical or different manner, by hydroxyl or carboxyl,

and/or the heterocycles mentioned under R^3 and R^4, formed together with the nitrogen atom, are optionally substituted by piperidinyl or pyrrolidinyl linked via N,

R^5 and R^6 are identical or different and represent hydrogen, straight-chain or branched alkyl having up to 6 carbon atoms, hydroxyl or straight-chain or branched alkoxy having up to 6 carbon atoms,

characterized in that compounds of the formula (II)

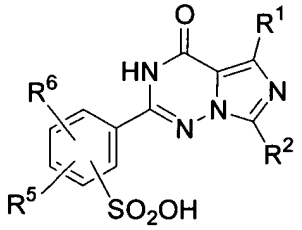


(II),

in which

R^1, R^2, R^5 and R^6 have the meanings indicated above,

are reacted with sulphuric acid to give compounds of the formula (III)

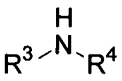


(III),

in which

R^1, R^2, R^5 and R^6 have the meanings indicated above,

and then with thionyl chloride and the product thus obtained is reacted in situ in an inert solvent with an amine of the formula (IV)



(IV),

in which

R^3 and R^4 have the meaning indicated above,

and, if appropriate, reacted to give the corresponding salts, hydrates or N-oxides.